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(71) Sökande Gambro Lundia AB, Lund SE
Applicant (s)

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REGISTRERINGSVERKET
SWEDEN

Postadress/Adress
Box 5055
S-102 42 STOCKHOLM

Telefon/Phone
+46 8 782 25 00
Vx 08-782 25 00

Telex
17978
PATOREG S

Telefax
+46 8 666 02 86
08-666 02 86

+46 40 260516

AWAPATENT AB

Kontor/Handläggare

Malmö/Dan Henriksson/JUN

GAMBRO LUNDIA AB

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METHOD FOR PREPARING A MEDICAL SOLUTION

2002-12-10

Huvudfaxen Kassar

Background of the Invention

The present invention relates to a method for preparing a medical solution, a solution used for preparing the medical solution, a container containing said solution, use of said solution for the manufacture of a medicament for peritoneal dialysis, and methods of treatment of patients with said medicament.

Background Art

N-acetylglucosamine (NAG) and glucosamine are biochemically classified as amino sugars. Amino sugars are formed in almost all cells from blood glucose through a series of biochemical reactions. Hyaluronan is a polymer composed of dimers containing N-acetylglucosamine and glucuronic acid. It has been shown that the function of the peritoneum as a dialysis membrane is better preserved in rats that have been chronically dialyzed with fluid supplemented with exogenous hyaluronan (see Wieczorowska K, Breborowicz A et al, Protective effect of hyaluronic acid against peritoneal injury, Perit Dial Int 1995; 15:81-3).

Breborowicz A, Kuzlan-Pawlaczyk M et al, The Effect of N-Acetyl-glucosamine as a Substrate for In Vitro Synthesis of Glycosaminoglycans by Human Peritoneal Mesothelial Cells and Fibroblasts, Advances in Peritoneal Dialysis, Vol. 14, 1998, teaches that NAG rapidly stimulates the production of hyaluronan and sulphated glycosaminoglycans by human peritoneal mesothelial cells and fibroblasts.

Wu G, Wieczorowska-Tobis K, et al, N-acetylglucosamine changes permeability of peritoneum during chronic peritoneal dialysis in rats, Perit Dial Int, Vol. 18, pp. 217-224 concludes that peritoneal dialysis with a dialysis solution supplemented with N-acetylglucosamin

causes accumulation of glucosaminoglycans in the peritoneal interstitium, resulting in a favourable change of the peritoneal permeability.

Due to these advantageous characteristics, NAG has been introduced as a component in peritoneal dialysis solutions replacing part or all of the glucose component with a view to obtaining a more biocompatible peritoneal dialysis solution (see WO97/06810).

Peritoneal dialysis is a method for exchanging solutes and water in capillary vessels of a patient's peritoneum with hypertonic solution, which is infused into the peritoneal cavity. The principle of this method is diffusion of solutes transferred according to the concentration gradient and water migration due to osmotic differences. This method has many advantages, e.g. that no special apparatus is commonly required. It gives less influence on the hemodynamics because extracorporeal circulation of the patient's blood is not necessary, and further the peritoneal dialysis is a continuous treatment and therefore more similar to the function of the kidneys.

Peritoneal dialysis is usually classified as continuous ambulatory peritoneal dialysis (CAPD), intermittent peritoneal dialysis (IPD), continuous cyclic peritoneal dialysis (CCPD) or automated peritoneal dialysis (APD).

In CAPD a catheter is permanently implanted in the abdominal wall of the patient and about 1.5 to 2.5 l of the dialysis fluid is normally introduced via the catheter into the peritoneal cavity. The peritoneal cavity is flooded with this fluid, left for an appropriate lapse of time and then drained. Removal of solutes and water takes place across the peritoneum which acts as a semipermeable membrane.

The dialysis fluid normally used for peritoneal dialysis is an aqueous solution comprising an osmotic agent such as glucose and the like, electrolytes such as sodi-

um, potassium, calcium, magnesium, and organic acid salts
such as sodium lactate, sodium bicarbonate, or sodium
pyruvate. The components of these peritoneal dialysis
fluids are selected to control the levels of electrolytes
5 or the acid-base equilibrium, to remove waste materials
and to efficiently carry out ultrafiltration.

It is known to pack medical solutions in multicom-
partment bags from e.g. WO 99/27885 (Gambro Lundia AB),
in which different solutes may be kept in separate com-
10 partments of the bag with a view to, inter alia, regulat-
ing the concentration of active ingredients in the fin-
ally prepared solution.

Medical fluids are normally sterilised by heat.
Medical authorities in many countries require sterilisa-
15 tion after final packaging of the medical product. It is
therefore often not possible to sterile filter the solu-
tion.

However, a problem with the formation of undesired
cytotoxic products during heat sterilisation and storage
20 exists for a variety of medical solutions, inter alia
within the dialysis area, e.g. for peritoneal dialysis
solutions. It is known e.g. from EP-B1-0 668 785 (Gambro
Lundia AB) to reduce the amount of toxic degradation pro-
ducts from glucose or glucose-like compounds in a medical
25 solution.

It has now been found that also amino sugars, e.g.
NAG, in conventional medical solutions exhibit an in-
creased cytotoxicity after heat sterilisation. This cyto-
toxicity depends on the formation of toxic degradation
30 products from said amino sugars. In contact to glucose,
none of the known glucose degradation products has been
found in heat sterilised NAG solutions. This fact has not
been known previously and forms the basis for the present
invention.

35 NAG and other amino sugars have a major difference
from glucose and glucose-like compounds by having one
amino group and possibly an acetyl group coupled to the

glucose ring. Regarding the degradation process, it has been found that the pH of a NAG solution increases during sterilisation while in the case of glucose it decreases during sterilisation. This indicates that NAG is, in contrast to glucose, degraded by a hydrolysis that forms acetate, which increases the pH.

Thus, there is a need to solve the above defined problem and to provide a medical solution containing amino sugars, in particular NAG, and derivatives thereof and at the same time having the ability to be heat sterilised without the formation of the above-mentioned cytotoxic products.

Summary of the Invention

The object of the present invention is to solve the above-mentioned problem.

According to the present invention this object is achieved by an improved method for preparing a medical solution, preferably a peritoneal dialysis solution, comprising the steps of:

a) providing a solution comprising one or more acetylated or deacetylated amino sugars in at least one compartment of a container, said solution having a pH of 2.0-5.0, and

b) sterilisation of said at least one compartment and the contents therein.

Further, the present invention relates to the solution used for preparing the medical solution, and to a container containing said solution.

The present invention also relates to use of said solution for the manufacture of a medicament for peritoneal dialysis.

In another aspect the present invention relates to a method of performing peritoneal dialysis, wherein said method comprises the introduction of said medicament for peritoneal dialysis into the peritoneal cavity of a patient.

In a further aspect the present invention relates to a method of treating a patient suffering from renal failure, said method comprising the introduction of said medicament for peritoneal dialysis into the peritoneal cavity of a patient.

In still another aspect the present invention relates to a method of reducing complications associated with peritoneal dialysis, said method comprising the introduction of said medicament for peritoneal dialysis into the peritoneal cavity of a patient.

Further disclosure of the objects, problems, solutions and features of the present invention will be apparent from the following detailed description of the invention with reference to the drawings and the appended claims.

Brief Description of the Drawings

Fig. 1 is a graph showing the relationship between pH and inhibition of cell growth (ICG) in a solution containing 1.5% NAG.

Fig. 2 is a bar diagram showing the effect of increased NAG concentration of 1.5% and 30% for three different pH values.

Fig. 3 is a graph showing the fluorescence of heat sterilised NAG containing solutions at different pH values.

Detailed Description of Preferred Embodiments

The present invention is a development of the above mentioned teachings and relates to a method for preparing a sterile medical solution, preferably a solution for peritoneal dialysis.

Experiments in which measurements of the percentage of inhibition of cell growth (ICG) and fluorescence have been made at varying pH values and NAG concentrations during sterilisation are illustrated in Figs 1-3. The results of the experiments imply that the pH of the amino sugar containing solution should be decreased from the neutral level and that, in a preferred embodiment of the

invention, also the concentration of the amino sugar/-sugars should be optimised.

A simple, reliable and known way to study the cytotoxicity of substances or of medical fluids is to test proliferation as in vitro inhibition of cell growth (ICG) in cultured cells. Another method to get a rough estimation of the amount of amino sugars that is rearranged is measurement of the fluorescence.

More precisely, from the graph in Fig. 1 it can be seen that the inhibition of cell growth reaches a minimum with sterilisation around pH 2.5-3.5 in a solution containing 1.5% NAG. This implies that a NAG containing solution sterilised at an optimal pH around 2.5-3.5 from an in vitro toxicological point of view is more compatible for humans than solutions giving a higher percentage of inhibited growth when sterilised at higher or lower pH values.

The bar diagram in Fig. 2 illustrates the effect of increased NAG concentration at different pH values during sterilisation. It can be seen that the percentage inhibition of cell growth is lower after sterilisation at a pH of 3.0 than at a pH of 5.5 and 7.2, and that the inhibition of cell growth is lower after a sterilisation at a NAG concentration of 30% than at a NAG concentration of 1.5% for all three pH values.

Fig. 3 shows the relationship between pH for a sterilised NAG containing solution and the fluorescence, measured at an excitation at 350 nm and an emission of 430 nm. The lowest fluorescence is seen at around pH 4, which corresponds well with the toxicity result shown in Fig. 1.

The sterilisation may include heat sterilisation or radiation sterilisation, but is preferably heat sterilisation effected in an autoclave at a temperature of at least 100°C, preferably at 121°C. The sterilisation time may vary depending on the sterilisation temperature, the

type of container and the contents therein to be sterilised.

The method according to the present invention is preferably effected for a multicompartment container as disclosed in WO 99/27885 (Gambro AB). In the present invention, such a container comprises at least one compartment containing a physiologically compatible pH adjusting and diluting solution as well as at least one compartment containing a solution comprising one or more acetylated or deacetylated amino sugars, in the following called amino sugar solution for simplicity. The amino sugar solution may be present in only one compartment. The solutions in the different compartments are heat sterilisable, and the whole container can be heat sterilised in an autoclave with the solutions in situ in said compartments. The solutions in the separated compartments, which are delimited from each other during the sterilisation and the subsequent storage, can be mixed after sterilisation to form a finally prepared sterile medical solution, preferably a solution for peritoneal dialysis. It may also be mixed with a sterilised pH adjusting and diluting solution in at least one other sterilised compartment of the container, thereby finally preparing the medical solution. Such a medical solution may be stored after sterilisation up to longer periods of time before mixing with the sterilised pH adjusting and diluting solution. The sterilisation can however also be effected for separated interconnectable containers comprising the solutions to sterilise and provided with connection means with sterile connecting valves for sterile connection. The separated containers can also be connected already during manufacture by means of a breakable seal, for example a conventional breakpin.

According to the invention, the pH of the amino
35 sugar containing solution is 2.0-5.0. In one preferred
embodiment of the invention the pH of the amino sugar
containing solution is preferably 2.5-3.5, most prefer-

ably 3.0, so that the formation of cytotoxic substances during the sterilisation step is substantially prevented.

In another embodiment of the invention the amino sugar in the amino sugar containing solution having a pH
5 of 2.0-5.0 in one or more of said compartments is present in a concentration of 15-40% by weight, preferably 20-40% by weight, most preferably at least 30% by weight, with the basis on the solution in each of said compartments, e.g. 15, 20, 25, 30, 35, and 40% by weight. Preferably,
10 said amino sugar is N-acetylglucosamine (NAG).

The upper limit for the concentration for each amino sugar in the solution is determined by its solubility therein. The compartment comprising the amino sugar containing solution may also contain any organic acid or
15 other pH stabilising agent in order to further stabilise the pH during sterilisation. The solutions of the different compartments have such respective pH values, concentrations and volumes that the finally prepared medical solution after mixing the solutions of the compartments
20 has a pH that is substantially neutral, i.e. a pH between 6.0 and 8.0, preferably about 7.4, and an amino sugar concentration between 0.2 and 15.0%, preferably 0.5-6.0%, e.g. 0.5-2.0% by weight, with the basis on the finally prepared solution.

The volume of each compartment, as well as the proportion between the compartments, is in practice not critical. Each compartment volume depends on the volume of constituent to be present therein. In the most preferred embodiment, the compartment which accommodates the
30 pH adjusting and diluting solution is larger than the compartment which accommodates the amino sugar containing solution and is also the compartment in which the solution/solutions from the other compartments is/are mixed with the pH adjusting and diluting solution.

In a preferred embodiment the medicament to be prepared is a peritoneal dialysis solution containing N-acetylglucoseamine and having a pH of 7.4.

The term "amino sugar containing solution" used herein means a solution comprising one or more acetylated or deacetylated amino sugars involved in the present invention chosen from N-acetylglucosamine (NAG), galactos-
5 amine, N-acetylgalactosamine, mannosamine, and N-acetylmannosamine in the form of monomers, oligomers and/or polymers thereof including chitin, and human glucosaminoglycans, as well as derivatives thereof. The most preferred amino sugar is N-acetylglucosamine (NAG). Thus,
10 the acetylated or deacetylated amino sugars may be represented by only one of the amino sugars listed or by a combination thereof as well as derivatives thereof.

The term "derivatives thereof" used herein means derivatives of said amino sugars having the same or essentially the same ability to form cytotoxic degradation
15 products during sterilisation.

The term "pH adjusting and diluting solution" used herein means a solution to be mixed with, e.g. acting as a receiving medium for, the amino sugar containing solution and at the same time a solution adjusting the pH of
20 the solution after mixing with the amino sugar containing solution to essentially neutral, i.e. with a pH between for example 6.0 and 8.0, preferably about 7.4.

The term "low levels of cytotoxic degradation products" used herein means that the amount of degradation
25 products from the amino sugars is so low in the medical solution prepared according to the present invention that it is not more toxic to cultured cells than dialysis solutions according to prior art.

30 The pH adjusting and diluting solution in the preferred embodiment contains pH adjusting agents, such as salts of inorganic acids, organic acids, alkalic substances etc. in a pharmaceutically stable range. Inorganic acids include hydrochloride acid etc., organic
35 acids include lactic acid, malic acid, acetic acid, succinic acid, maleic acid, pyruvic acid, citric acid, gluconic acid, etc., and alkalic substances include sodi-

um hydrate, sodium bicarbonate etc. Also, various amino acids can be used as a pH adjusting agent.

After sterilisation the amino sugar containing solution is finally prepared for use by mixing it with the pH adjusting and diluting solution, optionally with solutions in other compartments of the container. The medical solution, preferably a peritoneal dialysis solution, thus obtained may also comprise different electrolyte ions, e.g. sodium, potassium, calcium, magnesium, chloride, lactate, and bicarbonate ions, in concentrations which are biocompatible and substantially isotonic. The electrolytes may be originally present in the pH adjusting and diluting solution, the amino sugar containing solution and/or another solution in one or more other compartments of the container, depending on their compatibility during sterilisation and storage, normally in the form of pharmaceutically acceptable salts. The amount of cations in a peritoneal dialysis solution ready for use is generally 110 to 140 mEq/ml of sodium ions, 0 to 0.05 mEq/l of potassium ions, 0 to 3 mEq/l of magnesium ions and 0 to 6 mEq/l of calcium ions. Preferably the amount of chloride ions is 80 to 144 mEq/l.

The peritoneal dialysis solution as a preferred embodiment of the medical solution according to the present invention may also comprise other physiologically compatible constituents, e.g. further osmotic agents, such as carbohydrates, preferably glucose, proteins and peptides, preferably albumin, as well as antioxidants, such as bisulphite.

The peritoneal dialysis solution of the present invention described above is applicable not only to continuous ambulatory peritoneal dialysis (CAPD) but also to intermittent peritoneal dialysis (IPD), continuous cyclic peritoneal dialysis (CCPD), and automated peritoneal dialysis (APD). Moreover, it contains low levels of cytotoxic degradation products from amino sugars.

The present invention also relates to a solution as such having the above defined characteristics.

The present invention also relates to a container containing the amino sugar containing solution in at least one compartment, wherein said solution has been sterilised and contains low levels of cytotoxic degradation products.

Further, the present invention relates to use of the solution according to the present invention for the manufacture of a medicament for peritoneal dialysis, wherein it is mixed with a sterilised pH adjusting and diluting solution.

As stated above, the present invention also relates to a method of performing peritoneal dialysis, wherein said method comprises the introduction of said medicament for peritoneal dialysis into the peritoneal cavity of a patient.

The present invention also relates to a method of treating a patient suffering from renal failure, said method comprising the introduction of said medicament for peritoneal dialysis into the peritoneal cavity of a patient, e.g. a patient suffering from renal failure in combination with diabetes, obesity and/or hyperlipidemia.

Further, the present invention relates to a method of reducing complications associated with peritoneal dialysis, said method comprising the introduction of said medicament for peritoneal dialysis into the peritoneal cavity of a patient, wherein said complications associated with peritoneal dialysis in a preferred embodiment consist of:

- a) morphologic and functional deteriorations of the peritoneal membrane;
- b) peritonitis;
- c) adverse metabolic consequences and related cardiovascular diseases;
- d) protein malnutrition;
- e) hyperglycemia;

f) hyperinsulinemia;

g) hypertension; and any combination thereof.

In order to illustrate different embodiments of the present invention, containers having different compartment constructions containing the constituents for the preparation of a peritoneal dialysis solution will be described in the Examples below, as well as the composition of the solutions in each compartment. In the examples, N-acetylglucosamine (NAG) is used as amino sugar, either in one or two compartments of the container. The pH for the NAG containing solutions in each compartment varied between 2.0 and 5.0 before mixing and between 6.0 and 8.0 in the finally prepared medical solution.

Examples

Example 1

15	Compartment A	Volume	100 ml
		Sodium	0-140 mM
		NAG	300 g/l
20	Compartment B	Volume	180 ml
		Sodium	0-140 mM
		NAG	300 g/l
25	Compartment C	Volume	1900 ml
		Sodium	0-140 mM
		Lactate	40 mM
		Magnesium	0.25-0.75 mM
		Calcium	0.9-2.0 mM

Final composition when the contents of compartments A+C are mixed:

30	Volume	2000 ml
	Sodium	0-140 mM
	NAG	15 g/l
	Lactate	38 mM
	Magnesium	0.24-0.71 mM
	Calcium	0.85-1.9 mM

35 Final composition when the contents of compartments B+C are mixed:

Volume	2080 ml
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Huvudföreläsningen

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	Sodium	0-140 mM
	NAG	26 g/l
	Lactate	36.5 mM
	Magnesium	0.22-0.68 mM
5	Calcium	0.82-1.8 mM

Final composition when the contents of compartments A+B+C
are mixed:

	Volume	2180 ml
	Sodium	0-140 mM
10	NAG	38.5 g/l
	Lactate	34.9 mM
	Magnesium	0.21-0.65 mM
	Calcium	0.78-1.7 mM

15 Example 2

	Compartment A	Volume	100 ml
		Sodium	0-140 mM
		NAG	300 g/l
	Compartment B	Volume	180 ml
20		Sodium	0-140 mM
		Glucose	500 g/l
	Compartment C	Volume	1900 ml
		Sodium	0-140 mM
		Lactate	40 mM
25		Magnesium	0.25-0.75 mM
		Calcium	0.9-2.0 mM

Final composition when the contents of compartments A+C
are mixed:

	Volume	2000 ml
30	Sodium	0-140 mM
	NAG	15 g/l
	Glucose	0 g/l
	Lactate	38 mM
	Magnesium	0.24-0.71 mM
35	Calcium	0.86-1.9 mM

Final composition when the contents of compartments A+B+C
are mixed:

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Huvudutredning Krossen

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5	Volume	2100 ml
	Sodium	0-140 mM
	NAG	14.3 g/l
	Glucose	23.8 g/l
	Lactate	36 mM
	Magnesium	0.23-0.68 mM
	Calcium	0.81-1.8 mM

Example 3

10	Compartment A	Volume	60 ml
		Sodium	0-140 mM
		NAG	165 g/l
		Glucose	330 g/l
15	Compartment B	Volume	100 ml
		Sodium	0-140 mM
		NAG	165 g/l
		Glucose	330 g/l
20	Compartment C	Volume	1900 ml
		Sodium	0-140 mM
		Lactate	40 mM
		Magnesium	0.25-0.75 mM
		Calcium	0.9-2.0 mM

Final composition when the contents of compartments A+C are mixed:

25	Volume	1960 ml
	Sodium	0-140 mM
	NAG	5.1 g/l
	Glucose	10.1 g/l
	Lactate	38.8 mM
30	Magnesium	0.24-0.73 mM
	Calcium	0.87-1.9 mM

Final composition when the contents of compartments B+C are mixed:

35	Volume	2000 ml
	Sodium	0-140 mM
	NAG	8.25 g/l
	Glucose	16.5 g/l
	Lactate	38.0 mM

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Huvudföreläsningen

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Magnesium 0.24-0.71 mM

Calcium 0.86-1.9 mM

Final composition when the contents of compartments A+B+C
are mixed:

5

Volume 2060 ml

Sodium 0-140 mM

NAG 12.8 g/l

Glucose 25.6 g/l

Lactate 37 mM

10

Magnesium 0.23-0.69 mM

Calcium 0.83-1.8 mM

Example 4

Compartment A

Volume 1000 ml

NAG 10 g/l

15

Glucose 20 g/l

Magnesium 0.48-1.46 mM

Calcium 1.8-4.0 mM

Compartment B

Volume 1000 ml

Sodium 0-140 mM

20

Bicarbonate 165 g/l

Final composition when the contents of compartments A+B
are mixed:

Volume 2000 ml

Sodium 0-140 mM

25

NAG 5 g/l

Glucose 10 g/l

Bicarbonate 37.5 mM

Lactate 2.5 mM

30

Magnesium 0.24-0.73 mM

Calcium 0.9-2.0 mM

The invention has been described above with reference to preferred embodiments of the invention. A skilled person will recognise that further combinations are possible. Modifications which are apparent to a skilled person are intended to be incorporated within the

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Huvudfrågan: Känslan

scope of the invention, which is limited only by the
appended claims.

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CLAIMS

1. A method for preparing a medical solution, comprising the steps of:

5 a) providing a solution comprising one or more acetylated or deacetylated amino sugars in at least one compartment of a container, said solution having a pH of 2.0-5.0, and

10 b) sterilisation of said at least one compartment and the contents therein.

2. The method according to claim 1, wherein the pH is 2.5-3.5, preferably 3.0.

3. The method according to claim 1, wherein said one or more acetylated or deacetylated amino sugar/sugars
15 is/are chosen from N-acetylglucosamine (NAG), galactosamine, N-acetylgalactosamine, mannosamine, and N-acetylmannosamine in the form of monomers, oligomers and/or polymers thereof including chitin, and human glucose-aminoglycans, as well as derivatives thereof.

20 4. Method according to any one of the previous claims, wherein said one or more acetylated or deacetylated amino sugar/sugars is/are present in a concentration of 15-40% by weight, preferably 20-40% by weight, most preferably at least 30% by weight, with the basis of the
25 solution in said at least one compartment.

5. The method according to any one of the previous claims, wherein said one or more acetylated or deacetylated amino sugar/sugars is N-acetylglucosamine (NAG).

30 6. The method according to any one of the preceding claims, wherein the sterilisation is heat sterilisation at a temperature of at least 100°C, preferably at 121°C.

7. The method according to any one of the preceding claims, wherein each compartment of the container is delimited from the other/others during the sterilisation,
35 and wherein the sterilised solution containing one or more acetylated or deacetylated amino sugars is/are mixed with a sterilised pH adjusting and diluting solution in

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at least one other sterilised compartment of the container, thereby finally preparing the medical solution.

8. The method according to claim 7, wherein the pH in the finally prepared medical solution is 6.0-8.0, preferably 7.4.

9. The method according to claim 7 or 8, wherein the concentration of acetylated or deacetylated amino sugar/sugars in the finally prepared solution is/are 0.2-15.0% by weight, preferably 0.5-6.0% by weight.

10. The method according to any one of the preceding claims, wherein physiologically compatible constituents in the form of carbohydrates, preferably glucose, proteins, peptides, and antioxidants are present in one or more of said compartments.

11. The method according to any one of the preceding claims, wherein the medical solution prepared is a peritoneal dialysis solution.

12. A solution comprising one or more acetylated or deacetylated amino sugar/sugars and having a pH of 2.0-5.0, preferably 2.5-3.5, most preferably 3.0, wherein said solution is sterilised and contains low levels of cytotoxic degradation products.

13. The solution according to claim 12, wherein said one or more acetylated or deacetylated amino sugar/sugars is/are present in a concentration of 15-40% by weight, preferably 20-40% by weight, most preferably at least 30% by weight.

14. The solution according to any one of claims 12 and 13, wherein the acetylated or deacetylated amino sugar/sugars is/are as defined in claim 3, and preferably is N-acetylglucosamine.

15. A container comprising at least one compartment containing a solution according to any one of claims 12-14.

16. Use of a solution according to any one of claims 12-14 for the manufacture of a medicament for peritoneal

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Hovud'ensen Krossen

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dialysis, wherein it is mixed with a sterilised pH
adjusting and diluting solution.

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ABSTRACT

A method for preparing a medical solution, comprising the steps of a) providing a solution comprising one
 5 or more acetylated or deacetylated amino sugar/sugars in at least one compartment of a container at a pH from 2.5 to 5.0, and b) sterilisation of said at least one compartment and the contents therein, is disclosed, as well as a solution used for preparing the medical solution, a
 10 container containing said solution, use of said solution for the manufacture of a medicament for peritoneal dialysis, and methods of treatment of patients with said medicament.

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Election for publication: Fig 1.

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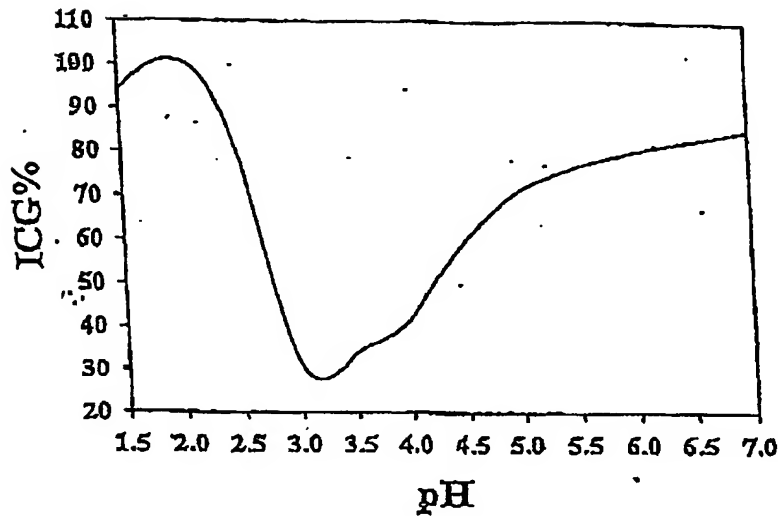


Figure 1.

Inhibition of cell growth, in a solution containing 1.5% NAG

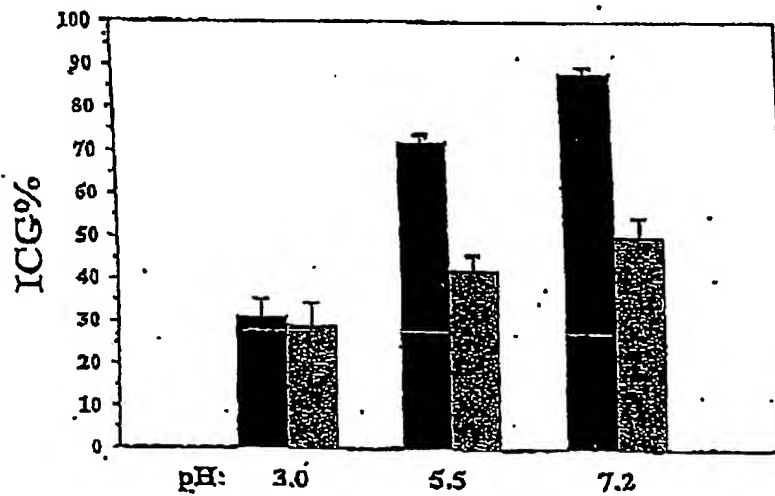


Figure 2.

The effect of increased NAG concentration from 1.5% (black bars) to 30% (grey bars) for three different pH values.

2/2

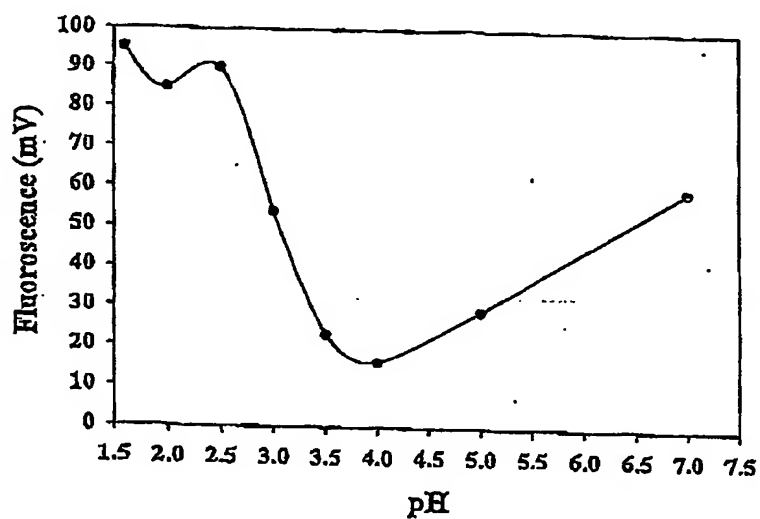


Figure 3.

Fluorescence of heat sterilised NAG containing solutions at different pH.

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